JUN 2 1 2005 W

PTO/SB/08B (07-05)
Approved for use through 6/30/2006. OMB 0651-0031
U.S. Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE
995, no persons are required to respond to a collection of information unless it contains a valid OMB control number.

Under the Paper 18 Reduction Act at 1995, no persons are required to respo

Complete if Known Substitute for form 1449/PTO 10/696,256 **Application Number** INFORMATION DISCLOSURE October 29, 2003 Filing Date STATEMENT BY APPLICANT Azrolan First Named Inventor (Use as many sheets as necessary) Group Art Unit 1614 Raymond J. Henley III **Examiner Name** AM-100302C1USA Attorney Docket Number of 1 Sheet

OTHER PRIOR ART-NONPATENT LITERATURE DOCUMENTS			
Examiner Initials*	Cite No.1	Include the name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item, (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue, number(s), publisher, city and/or country where published	T²
M	AAAA	Y. BOTTIGER, ET AL., Pharmacokinetic interaction between single oral doses of diltiazem and sirolimus, Clin. Pharmacol. Ther., January 2001, 69(1):32-40 (abstract).	
	AAAB	M. VALGIMIGLI, ET AL., Tirofiban and Sirolimus-Eluting Stent vs Abciximab and Bare-Metal Stent for Acute Myocardial Infarction, <i>JAMA</i> , May 4, 2005, Vol. 293, No. 17, pp. 2109-2117.	
	AAAC	J. MORRISETT, ET AL., Effects of sirolimus on plasma lipids, lipoprotein levels, and fatty acid metabolism in renal transplant patients, <i>Journal of Lipid Research</i> , Vol. 43, 2002, pp. 1170-1180.	
10	AAAD	Z. BERGER, ET AL., Rapamycin alleviates toxicity of different aggregrate-prone proteins, Human Molecular Genetics, 2006, Vol. 15, No. 3, pp. 433-442	
			_
			\vdash
······································			
			-
			<u> </u>

Examiner Signature

Date
Considered

4/28/06

* EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance

If you need assistance in completing the form, call 1-800-PTO-9199 (1-800-786-9199) and select option 2.

EXAMINER: Initial ir reference considered, whether or not clustum is in considered. Include copy of this form with next communication to applicant.

Applicant's unique citation designation number (optional). Applicant is to place a check mark here if English language Translation is attached. This collection of information is required by 37 CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is to flie (and by the USPTO to process) an application. Confidentiality is governed by 35 USC 122 and 37 CFR 1.14. this collection is estimated to take 2 hours to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, Washington, DC 20231. DO NOT SEND FEES OR COMPLETE FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. box 1450, Alexandria, VA 22313-1450.